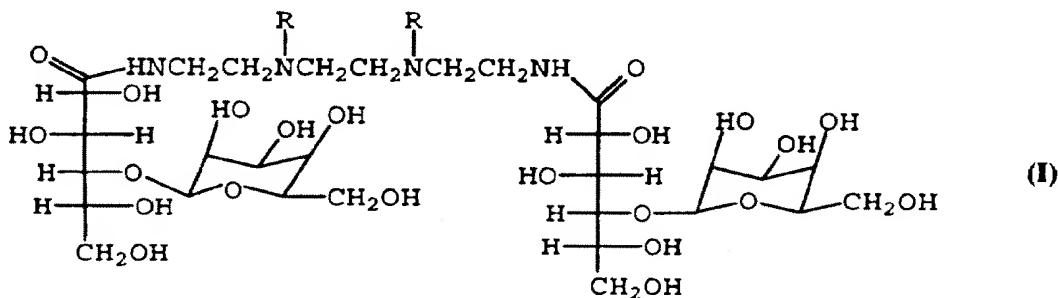




## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>6</sup> : <b>A61K 9/14, 9/51</b>		A1	(11) International Publication Number: <b>WO 96/24335</b>
			(43) International Publication Date: 15 August 1996 (15.08.96)
(21) International Application Number: PCT/US96/01439			(81) Designated States: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AZ, BY, KG, KZ, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).
(22) International Filing Date: 6 February 1996 (06.02.96)			
(30) Priority Data: 08/386,026 9 February 1995 (09.02.95) US 08/444,796 19 May 1995 (19.05.95) US			
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## (54) Title: SUGAR BASE SURFACTANT FOR NANOCRYSTALS



## (57) Abstract

Dispersible particles consisting essentially of a crystalline drug substance having a surface modifier having formula (I), wherein R is phenyl-(CH<sub>2</sub>)<sub>10</sub>CO or C<sub>12</sub>H<sub>25</sub>NHCO adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm, methods for the preparation of such particles and dispersions containing the particles. Pharmaceutical compositions containing the particles exhibit unexpected bioavailability and are useful in methods of treating mammals.

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SUGAR BASE SURFACTANT FOR NANOCRYSTALSFIELD OF THE INVENTION

This invention relates to drug particles, methods for the preparation thereof and 5 dispersions containing the particles. This invention further relates to the use of such particles in pharmaceutical compositions and methods of treating mammals.

BACKGROUND OF THE INVENTION

10 Bioavailability is the degree to which a drug becomes available to the target tissue after administration. Many factors can affect bioavailability including the dosage form and various properties, e.g., dissolution rate of the 15 drug. Poor bioavailability is a significant problem encountered in the development of pharmaceutical compositions, particularly those containing an active ingredient that is poorly soluble in water. Poorly water soluble drugs, 20 i.e., those having a solubility less than about 10 mg/ml, tend to be eliminated from the gastrointestinal tract before being absorbed into the circulation. Moreover, poorly water soluble drugs tend to be unsafe for intravenous 25 administration techniques, which are used primarily in conjunction with fully soluble drug substances.

It is known that the rate of 30 dissolution of a particulate drug can increase with increasing surface area, i.e., decreasing particle size. Consequently, methods of making finely divided drugs have been

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studied and efforts have been made to control the size and size range of drug particles in pharmaceutical compositions. For example, dry milling techniques have been used to reduce particle size and hence influence 5 drug absorption. However, in conventional dry milling, as discussed by Lachman, et al., The Theory and Practice of Industrial Pharmacy, Chapter 2, "Milling", p. 45, (1986), the limit of fineness is reached in the region of 100 microns (100,000 nm) when 10 material cakes on the milling chamber. Lachman, et al. note that wet grinding is beneficial in further reducing particle size, but that flocculation restricts the lower particle size limit to approximately 10 microns (10,000 nm). However, there tends to be a bias 15 in the pharmaceutical art against wet milling due to concerns associated with contamination. Commercial airjet milling techniques have provided particles ranging in average particle size from as low as about 1 to 50  $\mu\text{m}$  (1,000 - 50,000 nm).

20 Other techniques for preparing pharmaceutical compositions include loading drugs into liposomes or polymers, e.g., during emulsion polymerization. However, such techniques have problems and limitations. For example, a lipid soluble drug is often required in 25 preparing suitable liposomes. Further, unacceptably large amounts of the liposome or polymer are often required to prepare unit drug doses. Further still, techniques for preparing such pharmaceutical compositions tend to be complex. A principal technical 30 difficulty encountered with emulsion polymerization is the removal of contaminants, such as unreacted monomer or initiator, which can be toxic, at the end of the manufacturing process.

35 U. S. Patent 4,540,602 (Motoyama et al.) discloses a solid drug pulverized in an aqueous solution of a water-soluble high molecular substance using a wet

grinding machine. However, Motoyama et al. teach that as a result of such wet grinding, the drug is formed into finely divided particles ranging from 0.5  $\mu\text{m}$  (500 nm) or less to 5  $\mu\text{m}$  (5,000 nm) in diameter.

5 EPO 275,796 describes the production of colloidal dispersions comprising a substance in the form of spherical particles smaller than 500 nm. However, the method involves a precipitation effected by mixing a solution of the substance and a miscible  
10 non-solvent for the substance and results in the formation of non-crystalline nonoparticles. Furthermore, precipitation techniques for preparing particles tend to provide particles contaminated with solvents. Such solvents are often toxic and can be  
15 very difficult, if not impossible, to adequately remove to pharmaceutically acceptable levels to be practical.

U. S. Patent 4,107,288 describes particles in the size range from 10 to 1,000 nm containing a biologically or pharmacodynamically active material.  
20 However, the particles comprise a crosslinked matrix of macromolecules having the active material supported on or incorporated into the matrix.

U.S. Patent 5,145,684 describes unique nanoparticles comprising a crystal- drug substance  
25 having a surface modifier adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm, methods for the preparation of such particles and dispersions containing the particles. Pharmaceutical  
30 compositions containing the particles exhibit unexpected bioavailability and are useful in methods of treating mammals.

It would be desirable to provide even more stable  
35 dispersible drug particles in the submicron size range which can be readily prepared and which do not appreciably flocculate or agglomerate due to

interparticle attractive forces and do not require the presence of a crosslinked matrix. Moreover, it would be highly desirable to provide pharmaceutical compositions having enhanced bioavailability over those 5 described in U.S. Patent 5,145,684.

SUMMARY OF THE INVENTION

We have discovered stable, dispersible drug nanoparticles and a method for preparing such particles 10 by wet milling in the presence of grinding media in conjunction with a specific surface modifier. The particles can be formulated into pharmaceutical compositions exhibiting remarkably high bioavailability.

15 More specifically, in accordance with this invention, there are provided particles consisting essentially of a crystalline drug substance having a surface modifier adsorbed on the surface thereof in an amount sufficient to maintain an effective average 20 particle size of less than about 400 nm.

This invention also provides a stable dispersion consisting essentially of a liquid dispersion medium and the above-described particles dispersed therein.

In another embodiment of the invention, there is 25 provided a method of preparing the above-described particles comprising the steps of dispersing a drug substance in a liquid dispersion medium and applying mechanical means in the presence of grinding media to reduce the particle size of the drug substance to an 30 effective average particle size of less than about 400 nm. The particles can be reduced in size in the presence of a surface modifier. Alternatively, the particles can be contacted with the surface modifier after attrition.

35 In a particularly valuable and important embodiment of the invention, there is provided a

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pharmaceutical composition comprising the above-described particles and a pharmaceutically acceptable carrier therefor. Such pharmaceutical composition is useful in a method of treating mammals.

5 It is an advantageous feature that a wide variety of surface modified drug nanoparticles free of unacceptable contamination can be prepared in accordance with this invention.

10 It is another advantageous feature of this invention that there is provided a simple and convenient method for preparing drug nanoparticles by wet milling in conjunction with the surface modifier.

15 Another particularly advantageous feature of this invention is that pharmaceutical compositions are provided exhibiting unexpectedly high bioavailability.

20 Still another advantageous feature of this invention is that pharmaceutical compositions containing poorly water soluble drug substances are provided which are suitable for intravenous administration techniques.

Other advantageous features will become readily apparent upon reference to the following Description of Preferred Embodiments.

25 DESCRIPTION OF PREFERRED EMBODIMENTS

This invention is based on the discovery that drug particles having an extremely small effective average particle size can be prepared by wet milling in the presence of grinding media in conjunction with a specific surface modifier, and that such particles are stable and do not appreciably flocculate or agglomerate due to interparticle attractive forces and can be formulated into pharmaceutical compositions exhibiting unexpectedly high bioavailability. While the invention is described herein primarily in connection with its preferred utility, i.e., with respect to

nanoparticulate drug substances for use in pharmaceutical compositions, it is also believed to be useful in other applications such as the formulation of particulate cosmetic compositions and the preparation 5 of particulate dispersions for use in image and magnetic recording elements.

The particles of this invention comprise a drug substance. The drug substance exists as a discrete, crystalline phase. The crystalline phase differs from 10 a non-crystalline or amorphous phase which results from precipitation techniques, such as described in EPO 275,796 cited above.

The invention can be practiced with a wide variety of drug substances. The drug substance preferably is 15 present in an essentially pure form. The drug substance must be poorly soluble and dispersible in at least one liquid medium. By "poorly soluble" it is meant that the drug substance has a solubility in the liquid dispersion medium of less than about 10 mg/ml, 20 and preferably of less than about 1 mg/ml. A preferred liquid dispersion medium is water. However, the invention can be practiced with other liquid media in which a drug substance is poorly soluble and dispersible including, for example, aqueous salt 25 solutions, safflower oil and solvents such as ethanol, t-butanol, hexane and glycol. The pH of the aqueous dispersion media can be adjusted by techniques known in the art.

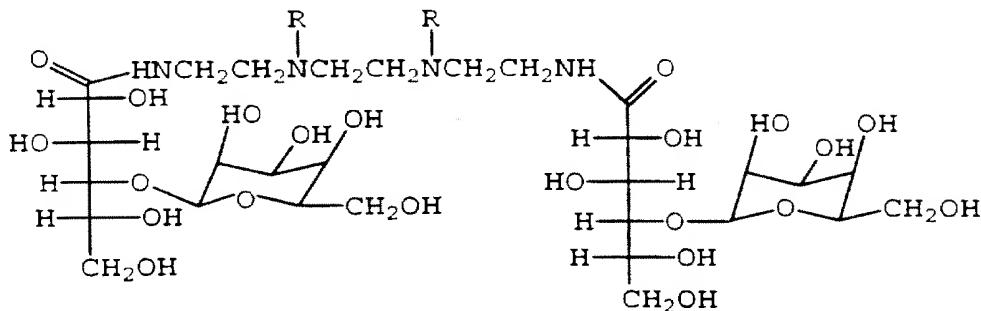
Suitable drug substances can be selected from a 30 variety of known classes of drugs including, for example, analgesics, anti-inflammatory agents, anthelmintics, anti-arrhythmic agents, antibiotics (including penicillins), anticoagulants, antidepressants, antidiabetic agents, antiepileptics, 35 antihistamines, antihypertensive agents, antimuscarinic agents, antimycobacterial agents, antineoplastic

agents, immunosuppressants, antithyroid agents, antiviral agents, anxiolytic sedatives (hypnotics and neuroleptics), astringents, beta-adrenoceptor blocking agents, blood products and substitutes, cardiac 5 inotropic agents, contrast media, corticosteroids, cough suppressants (expectorants and mucolytics), diagnostic agents, diagnostic imaging agents, diuretics, dopaminergics (antiparkinsonian agents), haemostatics, immunological agents, lipid regulating 10 agents, muscle relaxants, parasympathomimetics, parathyroid calcitonin and biphosphonates, prostaglandins, radio-pharmaceuticals, sex hormones (including steroids), anti-allergic agents, stimulants and anoretics, sympathomimetics, thyroid agents, 15 vasodilators and xanthines. Preferred drug substances include those intended for oral administration and intravenous administration. A description of these classes of drugs and a listing of species within each class can be found in Martindale, *The Extra 20 Pharmacopoeia*, Twenty-ninth Edition, The Pharmaceutical Press, London, 1989, the disclosure of which is hereby incorporated by reference in its entirety. The drug substances are commercially available and/or can be prepared by techniques known in the art.

25 Representative illustrative species of drug substances useful in the practice of this invention include:

30 17-pregno-2,4-dien-20-yne-[2,3-d]-isoxazol-17-ol  
(Danazol);  
5,17,-1'-(methylsulfonyl)-1'H-pregn-20-yne  
[3,2-c]-pyrazol-17-ol (Steroid A);  
piposulfam;  
piposulfan;  
35 camptothecin; and  
ethyl-3,5-diacetoamido-2,4,6-triiodobenzoate

In particularly preferred embodiments of the invention, the drug substance is a diagnostic agent surface modifier having the structure:



5

wherein R is  $\text{Ph}(\text{CH}_2)_{10}\text{CO}$  wherein Ph is phenyl or  $\text{C}_{12}\text{H}_{25}\text{NHCO}$  adsorbed on the surface thereof.

The particular surface modifier provides the nanodispersible particles with unexpectedly reduced mean particle size and limiting the particle size growth during terminal sterilization of the nanocrystal formulations.

The surface modifier of this invention is the discovery of Ian Newington and Katie Adams.

15 The surface modifier is adsorbed on the surface of the drug substance in an amount sufficient to maintain an effective average particle size of less than about 400 nm. The surface modifier does not chemically react with the drug substance or itself. Furthermore, the 20 individually adsorbed molecules of the surface modifier are essentially free of intermolecular crosslinkages.

As used herein, particle size refers to a number average particle size as measured by conventional particle size measuring techniques well known to those skilled in the art, such as sedimentation field flow fractionation, photon correlation spectroscopy, or disk centrifugation. By "an effective average particle size of less than about 400 nm" it is meant that at least 90% of the particles have a weight average particle

size of less than about 400 nm when measured by the above-noted techniques. In preferred embodiments of the invention, the effective average particle size is less than about 250 nm. In some embodiments of the 5 invention, an effective average particle size of less than about 100 nm has been achieved. With reference to the effective average particle size, it is preferred that at least 95% and, more preferably, at least 99% of the particles have a particle size less than the 10 effective average, e.g., 400 nm. In particularly preferred embodiments, essentially all of the particles have a size less than 400 nm. In some embodiments, essentially all of the particles have a size less than 250 nm.

15 The particles of this invention can be prepared in a method comprising the steps of dispersing a drug substance in a liquid dispersion medium and applying mechanical means in the presence of grinding media to reduce the particle size of the drug substance to an 20 effective average particle size of less than about 400 nm. The particles can be reduced in size in the presence of a surface modifier. Alternatively, the particles can be contacted with a surface modifier after attrition.

25 A general procedure for preparing the particles of this invention is set forth below. The drug substance selected is obtained commercially and/or prepared by techniques known in the art in a conventional coarse form. It is preferred, but not essential, that the 30 particle size of the coarse drug substance selected be less than about 100  $\mu\text{m}$  as determined by sieve analysis. If the coarse particle size of the drug substance is greater than about 100  $\mu\text{m}$ , then it is preferred that the particles of the drug substance be reduced in size 35 to less than 100  $\mu\text{m}$  using a conventional milling method such as airjet or fragmentation milling.

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The coarse drug substance selected can then be added to a liquid medium in which it is essentially insoluble to form a premix. The concentration of the drug substance in the liquid medium can vary from about 5 0.1 - 60%, and preferably is from 5 - 30% (w/w). It is preferred, but not essential, that the surface modifier be present in the premix. The concentration of the surface modifier can vary from about 0.1 to about 90%, and preferably is 1 - 75%, more preferably 20-60%, by 10 weight based on the total combined weight of the drug substance and surface modifier. The apparent viscosity of the premix suspension is preferably less than about 1000 centipoise

The premix can be used directly by subjecting it 15 to mechanical means to reduce the average particle size in the dispersion to less than 400 nm. It is preferred that the premix be used directly when a ball mill is used for attrition. Alternatively, the drug substance and, optionally, the surface modifier, can be dispersed 20 in the liquid medium using suitable agitation, e.g., a roller mill or a Cowles type mixer, until a homogeneous dispersion is observed in which there are no large agglomerates visible to the naked eye. It is preferred that the premix be subjected to such a premilling 25 dispersion step when a recirculating media mill is used for attrition.

The mechanical means applied to reduce the particle size of the drug substance conveniently can take the form of a dispersion mill. Suitable 30 dispersion mills include a ball mill, an attritor mill, a vibratory mill, and media mills such as a sand mill and a bead mill. A media mill is preferred due to the relatively shorter milling time required to provide the intended result, i.e., the desired reduction in 35 particle size. For media milling, the apparent viscosity of the premix preferably is from about 100 to

about 1000 centipoise. For ball milling, the apparent viscosity of the premix preferably is from about 1 up to about 100 centipoise. Such ranges tend to afford an optimal balance between efficient particle fragmentation and media erosion.

The grinding media for the particle size reduction step can be selected from rigid media preferably spherical or particulate in form having an average size less than about 3 mm and, more preferably, less than about 1 mm. Such media desirably can provide the particles of the invention with shorter processing times and impart less wear to the milling equipment. The selection of material for the grinding media is not believed to be critical. We have found that zirconium oxide, such as 95% ZrO stabilized with magnesia, zirconium silicate, and glass grinding media provide particles having levels of contamination which are believed to be acceptable for the preparation of pharmaceutical compositions. However, other media, such as stainless steel, titania, alumina, and 95% ZrO stabilized with yttrium, are expected to be useful. Preferred media have a density greater than about 3 g/cm<sup>3</sup>.

The attrition time can vary widely and depends primarily upon the particular mechanical means and processing conditions selected. For ball mills, processing times of up to five days or longer may be required. On the other hand, processing times of less than 1 day (residence times of one minute up to several hours) have provided the desired results using a high shear media mill.

The particles must be reduced in size at a temperature which does not significantly degrade the drug substance. Processing temperatures of less than about 30 - 40°C are ordinarily preferred. If desired, the processing equipment can be cooled with

conventional cooling equipment. The method is conveniently carried out under conditions of ambient temperature and at processing pressures which are safe and effective for the milling process. For example, 5 ambient processing pressures are typical of ball mills, attritor mills and vibratory mills. Processing pressures up to about 20 psi (1.4 kg/cm<sup>2</sup>) are typical of media milling.

10 The surface modifier, if it was not present in the premix, must be added to the dispersion after attrition in an amount as described for the premix above.

15 Thereafter, the dispersion can be mixed, e.g., by shaking vigorously. Optionally, the dispersion can be subjected to a sonication step, e.g., using an ultrasonic power supply. For example, the dispersion can be subjected to ultrasonic energy having a frequency of 20 - 80 kHz for a time of about 1 to 120 seconds.

20 The relative amount of drug substance and surface modifier can vary widely and the optimal amount of the surface modifier can depend, for example, upon the particular drug substance and surface modifier selected, the critical micelle concentration of the surface modifier if it forms micelles, etc. The 25 surface modifier preferably is present in an amount of about 0.1-10 mg per square meter surface area of the drug substance. The surface modifier can be present in an amount of 0.1-90%, preferably 20-60% by weight based on the total weight of the dry particle.

30 The resulting dispersion of this invention is stable and consists of the liquid dispersion medium and the above-described particles. The dispersion of surface modified drug nanoparticles can be spray coated onto sugar spheres or onto a pharmaceutical excipient 35 in a fluid-bed spray coater by techniques well known in the art.

Pharmaceutical compositions according to this invention include the particles described above and a pharmaceutically acceptable carrier therefor. Suitable pharmaceutically acceptable carriers are well known to those skilled in the art. These include non-toxic physiologically acceptable carriers, adjuvants or vehicles for parenteral injection, for oral administration in solid or liquid form, for rectal administration, and the like. A method of treating a mammal in accordance with this invention comprises the step of administering to the mammal in need of treatment an effective amount of the above-described pharmaceutical composition. The selected dosage level of the drug substance for treatment is effective to obtain a desired therapeutic response for a particular composition and method of administration. The selected dosage level therefore, depends upon the particular drug substance, the desired therapeutic effect, on the route of administration, on the desired duration of treatment and other factors. As noted, it is a particularly advantageous feature that the pharmaceutical compositions of this invention exhibit unexpectedly high bioavailability as illustrated in the examples which follow. Furthermore, it is contemplated that the drug particles of this invention provide more rapid onset of drug action and decreased gastrointestinal irritancy.

It is contemplated that the pharmaceutical compositions of this invention will be particularly useful in oral and parenteral, including intravenous, administration applications. It is expected that poorly water soluble drug substances, which prior to this invention, could not have been administered intravenously, may be administered safely in accordance with this invention. Additionally, drug substances which could not have been administered orally due to

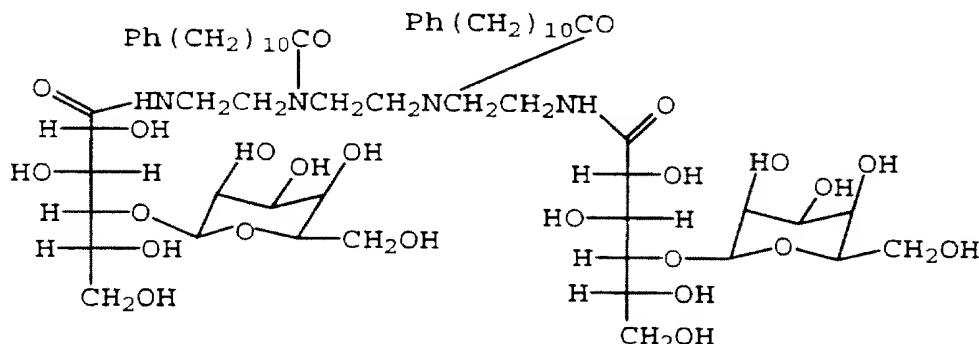
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poor bioavailability may be effectively administered in accordance with this invention.

The following Examples further illustrate the invention.

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**Example 1: Preparation of SA90HEA**

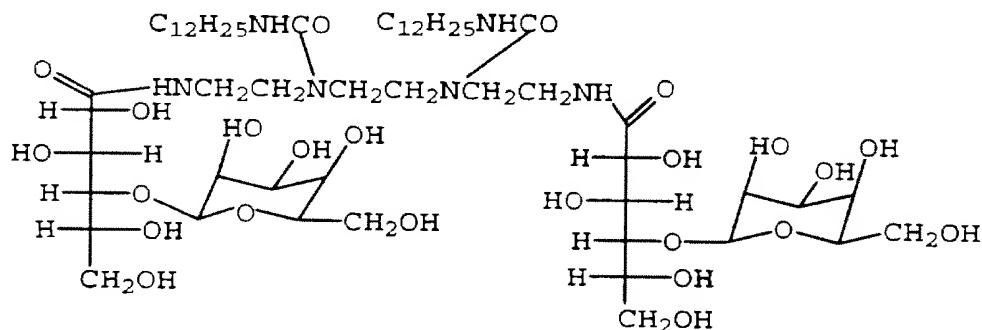


To a solution of lactobionic acid (17.92g, 50mmol, 2 equivs. Aldrich 97%) in DMF (Dimethylformamide, 100ml, dried over 4A° molecular sieves). The reaction mixture was stirred at 70°C under argon for 18 hours. The solution was cooled to 50°C. The resulting product was N, N10-triethylenetetramine-bislactobionamide. N4,N7-diphenylundecanoyl-N1,N10-triethylenetetramine-bislactobionamide-SA90HEA as prepared by dissolving phenylundecanoic acid (13.12g, 50mmol, 2 equivs. Eastman Chemicals 99%) in diethyl ether (40ml) and cooling in an ice-bath. Triethylamine (5.06g, 500mmol, 2 equivs, Prolabo) was added followed by ethyl chloroformate (5.53g, 51mmol, 2.04 equivs. Lancaster), the mixture was stirred well. After 30 mins. filtered into a solution of N1, N10-triethylenetetraminebislactobionamide (20.67g, 25mmol, 1 equiv) in DMF (100ml, dried over 4A° molecular sieves) and washed well with diethyl ether. The mixture was stirred at 50°C for 6 hours. After cooling to room temperature, the solvent was evaporated under reduced pressure to leave an orange crystalline material, which was washed with diethyl ether.

collected by Buchner filtration and briefly dried under high vacuum to leave an orange powder (30.72g, 93% yield). A portion of this material (13g) was then dissolved in distilled water (40ml) and stirred with 5 Amberlite IRA-420(OH) resin (BDH) at pH=10 for 40 mins. The resin was removed by buchner filtration and the product was obtained after freeze drying overnight which removed the water. SA90HEA was obtained as a fine pale yellow powder (7.47g, 57% yield).

10

Example 2: Preparation of SA90HEG



N4,N7-didodecylisocyano-N1,N10-triethylenetetramine-bislactobionamide - SA90HEG was 15 prepared by adding Dodecyl isocyanate (10.57g, 50mmol, 2 equivs. Eastman Chemicals 95%) to a solution of N1, N10-triethylenetetraminebislactobionamide (20.67g, 25mmol, 1 equiv) in DMF (100ml, dried over 4A° molecular sieves). The reaction mixture was heated at 20 50°C under argon for 7 hours. After cooling to room temperature, the solvent was evaporated under reduced pressure to leave a yellow crystalline material, which was washed with diethyl ether, collected by Buchner filtration and briefly dried under high vacuum to leave 25 a yellow powder (29.15g, 93% yield). A portion of this material (16g) was then dissolved in distilled water (50ml) and stirred with Amberlite IRA-420(OH) resin (BDH) at pH=10 for 40 mins. The resin was removed by Buchner filtration and the product was obtained after

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freeze drying overnight which removed the water.

SA90HEG was obtained as a fine cream powder (11.19g, 70% yield).

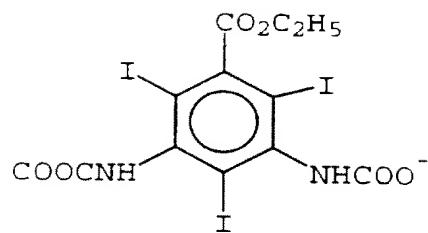
5 Examples 3-6

The following formulations were prepared at 15% diagnostic agent and 4% surfactant (w/v). A 6% stock solution was prepared by dissolving 600 mg of SA90HEA and SA90HEG in 10 ml deionized water. To each 15 ml 10 amber colored bottle, 7.5 ml of 6% stock surfactant solution and 0.994 ml deionized water were added. The sample bottle was sealed and placed on a roller mill running at 160 rpm for 7 days. At day 7, aliquot of samples were diluted 100 fold with deionized water for 15 particle size measurement by Photon Correlation Spectroscopy (Microtrac).

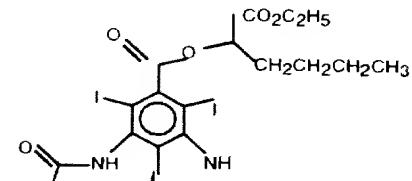
For terminal sterilization, 1 ml of nanocrystal formulation prepared above was pipetted into a 2 ml serum vial. After sealing with rubber septum and 20 aluminum cap, the samples were subjected to autoclave at 121°C for 20 min. On cooling to room temperature, aliquots of samples were retrieved from the vial and diluted 100 fold with deionized water for particle size measurement by Photon Correlation Spectroscopy 25 (Microtrac).

Example	Core	Surfactant	Mean Particle Size (nm)	
			Before Auto	After Auto
3	Compound A	SA90HEA	55	125
4	Compound A	SA90HEG	87	194
A	Compound A	F108	130	>500
B	Compound A	T908	241	>500
5	Compound B	SA90HEA	77	
6	Compound B	SA90HEG	71	

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Compound A = WIN 8883



Compound B = WIN 70146

5 Comparing to the commercially available copolymeric surfactants under identical milling and autoclave conditions, this data demonstrates that the sugar surfactants, SA90HEA and SA90HEG, resulted in unexpectedly reduced mean particle size and limited the  
10 particle size growth during terminal sterilization of the nanocrystal formulations.

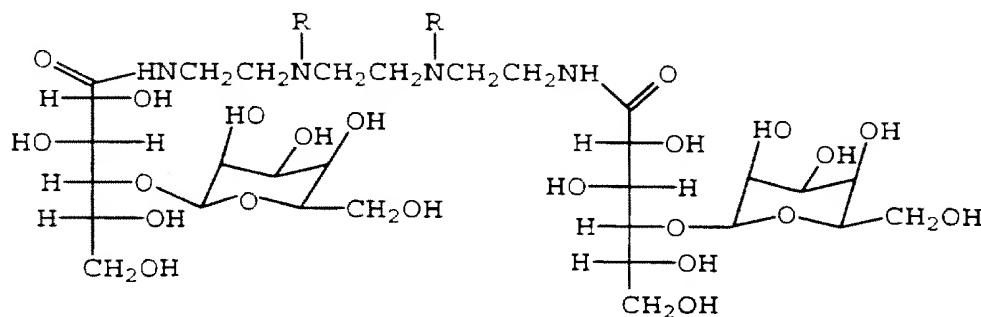
15 In addition, tail vein injection of a 4% solution of SA90HEA and SA90HEG at 30 ml/Kg was well tolerated by mice. SA90HEA and SA90HEG were tested in a smudge cell evaluation. Both SA90HEG and SA90HEA were devoid of smudge cell effect.

SA90HEA and SA90HEG also significantly inhibited the uptake of polystyrene particle by macrophages.

20 The invention has been described in detail with particular reference to preferred embodiments thereof, but it will be understood that variations and modifications can be effected within the spirit and scope of the invention.

What is claimed is:

1. Particles consisting essentially of a crystalline drug substance having a surface modifier 5 having the structure:



10 wherein R is  $\text{Ph}(\text{CH}_2)_{10}\text{CO}$  or  $\text{C}_{12}\text{H}_{25}\text{NHCO}$  adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm.

15 2. The particles of claim 1 having an effective average particle size of less than 250 nm.

3. The particles of claim 1 having an effective average particle size of less than 100 nm.

20 4. The particles of claim 1 wherein said drug substance is selected from analgesics, anti-inflammatory agents, anthelmintics, anti-arrhythmic agents, antibiotics, anticoagulants, antidepressants, antidiabetic agents, antiepileptics, antihistamines, 25 antihypertensive agents, antimuscarinic agents, antimycobacterial agents, antineoplastic agents, immunosuppressants, antithyroid agents, antiviral agents, anxiolytic sedatives, astringents, beta-adrenoceptor blocking agents, contrast media, corticosteroids, cough suppressants, diagnostic agents, diagnostic imaging agents, diuretics, dopaminergics, haemostatics, immunochemical agents, lipid regulating

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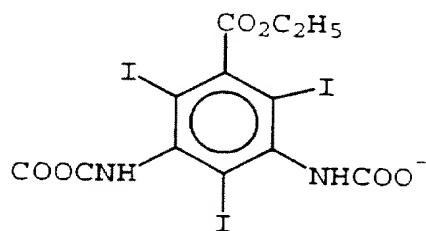
agents, muscle relaxants, parasympathomimetics, parathyroid calcitonin, prostaglandins, radio-pharmaceuticals, sex hormones, anti-allergic agents, stimulants, sympathomimetics, thyroid agents,

5 vasodilators and xanthines.

5. The particles of claim 1 wherein the drug substance is a diagnostic agent.

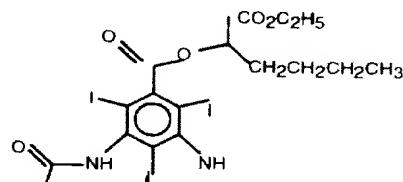
6. The particles of claim 5 wherein the drug substance is

10

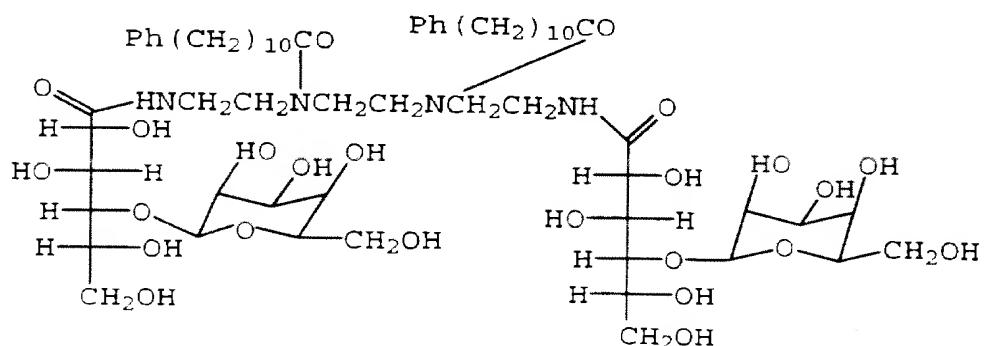


7. The particles of claim 5 wherein the drug substance is

15

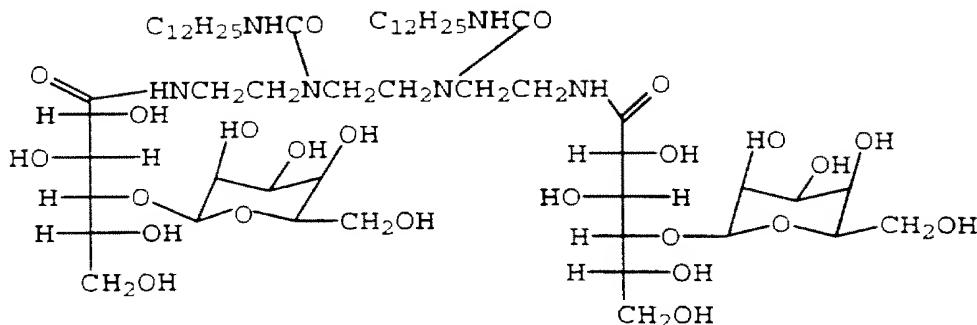


8. The particles of claim 1 wherein the surface modifier is



9. The particles of claim 1 wherein the surface 20 modifier is

-20-



10. Particles consisting essentially of a discrete phase of 5,17,-1'-(methylsulfonyl)-1'H-pregn-20-yne-[3,2-c]-pyrazol-17-ol having an ethylene oxide propylene-oxide block copolymer adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm.

11. A stable dispersion consisting essentially of 10 a liquid dispersion medium and the particles of claim 1.

12. The dispersion of claim 10 wherein said dispersion medium is water.

13. The dispersion of claim 10 wherein said dispersion medium is selected from the group consisting of safflower oil, ethanol, t-butanol, hexane and glycol.

14. A pharmaceutical composition comprising the particles of claim 1 and a pharmaceutically acceptable 20 carrier therefor.

15. A method of treating a mammal comprising the step of administering to the mammal an effective amount of the pharmaceutical composition of claim 13.

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 96/01439

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 6 A61K9/14 A61K9/51

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, Y	EP,A,0 688 781 (EASTMAN KODAK COMPANY) 27 December 1995 see the whole document ---	1-9, 11-15
Y	EP,A,0 603 944 (KODAK LIMITED) 29 June 1994 see claims 1-12 ---	1-9, 11-15
Y	EP,A,0 498 482 (STERLING WINTHROP INC.) 12 August 1992 see page 3, line 13 - line 24 see page 5, line 1 - page 8, line 38 -----	1-9, 11-15

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

**\* Special categories of cited documents :**

- \*'A' document defining the general state of the art which is not considered to be of particular relevance
- \*'E' earlier document but published on or after the international filing date
- \*'L' document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*'O' document referring to an oral disclosure, use, exhibition or other means
- \*'P' document published prior to the international filing date but later than the priority date claimed

- \*'T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- \*'X' document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- \*'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- \*'&' document member of the same patent family

Date of the actual completion of the international search

8 May 1996

Date of mailing of the international search report

17.07.96

Name and mailing address of the ISA

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Tzschoppe, D

**INTERNATIONAL SEARCH REPORT**

International Application No  
PCT/US 96/01439

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
EP-A-0688781	27-12-95	JP-A- US-A-	8041094 5529897	13-02-96 25-06-96
EP-A-0603944	29-06-94	JP-A- US-A-	6228069 5457023	16-08-94 10-10-95
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## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 96/01439

### Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
  
2.  Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
  
3.  Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

### Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. claims 1-9, 11-15

2. claim 10

For further information, please see form PCT/ISA/206 dated 30.05.96.

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
  
2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
  
3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-9, 11-15

#### Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

